



Food and Drug Administration  
Rockville, MD 20857  
Re: CEREBYX

Docket No. 96E-0442

JAN 21 1997

# 18

Stephen G. Kunin  
Deputy Assistant Commissioner for  
Patent Policy and Projects  
U.S. Patent and Trademark Office  
Box Pat. Ext.  
Assistant Commissioner for Patents  
Washington, D.C. 20231

Dear Mr. Kunin:

This is in regard to the application for patent term extension for U.S. Patent No. 4,260,769 filed by Warner-Lambert Company under 35 U.S.C. § 156. The human drug product claimed by the patent is CEREBYX (fosphenytoin sodium), which was assigned New Drug Application (NDA) No. 20-450.

A review of the Food and Drug Administration's official records indicates that this product was subject to a regulatory review period before its commercial marketing or use, as required under 35 U.S.C. § 156(a)(4). Our records also indicate that it represents the first permitted commercial marketing or use of the product, as defined under 35 U.S.C. § 156(f)(1), and interpreted by the courts in Glaxo Operations UK Ltd. v. Quigg, 706 F. Supp. 1224 (E.D. Va. 1989), aff'd, 894 F.2d 392 (Fed. Cir. 1990).

The NDA was approved on August 5, 1996, which makes the submission of the patent term extension application on September 30, 1996, timely within the meaning of 35 U.S.C. § 156(d)(1).

Should you conclude that the subject patent is eligible for patent term extension, please advise us accordingly. As required by 35 U.S.C. § 156(d)(2)(A) we will then determine the applicable regulatory review period, publish the determination in the Federal Register, and notify you of our determination.

Please let me know if we can be of further assistance.

Sincerely,

Ronald L. Wilson, Director  
Health Assessment Policy Staff  
Office of Health Affairs

cc: Warner-Lambert Company / Patent Department  
Todd M. Crissey  
2800 Plymouth Road  
Ann Arbor, MI 48105